REMARKS

Presented herein are new claims 8-14 that correspond, respectively, to originally presented claims 1-7, before amendments made under Article 34, except that all multiple dependency has been eliminated. Claims 15 and 16 are like claims 10 and 11 but depend on claim 9 rather than claim 8.

A substitute specification that employs more-idiomatic English is submitted herewith. A clean version and a version showing the changes are both submitted. The substitute specification contains no new matter. The amendments herein to the specification are the Article 34 specification amendments. They describe in words examples originally presented in Table 1.

An early examination on the merits is respectfully requested.

The Commissioner is hereby authorized to charge any deficiency in fees or to credit any overpayment in fees to Attorney's Deposit Account No. 50-0562.

Respectfully submitted,

Date: 4-25-05

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SPECIFICATION

POLYPROPYLENE FIBER FOR CEMENT REINFORCEMENT, MOLDED CEMENT MADE
WITH THE FIBER, METHOD OF CONSTRUCTING CONCRETE STRUCTURE, AND
METHOD OF CONCRETE SPRAYING

FIELD OF THE INVENTION

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The present invention relates to polypropylene resin fibers for cement reinforcement excellent in adhesive property to cement matrixes and in reinforcing mortars, concrete and the like. In addition, the invention relates to fiber-reinforced molded cement made with the polypropylene resin fibers, a method of constructing concrete structure therewith, and a method of concrete spraying therewith.

BACKGROUND OF THE INVENTION

Conventionally, cement moldings, external walls of architectures, inner walls of tunnels and hillside slopes, etc. are constructed with mortars, concrete and the like. However, these moldings have are comparatively fragilelarge fragility. Therefore, unless properties the property such as the reinforced strength, flexural proof stress, flexural toughness and impact strength thereof areis enough, a water leaks and an external-wall peeling or dropping accidents happen due to cracks of wall surfaces. Therefore, recently, synthetic resin fibers such as steel fibers, polyvinylalcohol resin, polyolefine resin, polyacrylonitrile resin and polyamide resin, have been used as reinforcement for the cement moldings. (For example, see Japanese Patent Laid-open Publication No. 303877 of 1988, Japanese Patent Laid-open Publication No. 218220 of 1996, Japanese Patent Laid-open Publication No. 86984 of 1997 and Japanese Patent Publication No. 40786 of 1989.)

However, as for the concrete in which the steel fibers are entrapped, since the specific gravity of steel fibers is as heavy as 7.8, materials including them is are difficult to carrybe carried or be entrapped. In addition, as for shotcrete, when the concrete is sprayed, the splashed and dropped, steel fibers are sometimes stepped on, causing to cause workers to be injured. Moreover, there is

another problem as the steel fibers get rusty.

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The eConcrete in which the polyvinylalcohol resin fibers are entrapped has the following problems. The fibers themselves absorbhave water absorption and hydrolyze at a high temperature under an alkali condition. Conversely, the concrete without the fibers is remarkably apt to decrease in the slump flow, and therefore, athe per unit amount of water must be increased to maintain enough slump flow for the spraying.

Recently, the cement moldings are often have been cured with an autoclave to improve dimensional stability or to shorten a curing time. However, this autoclave curing has a problem that such as the fibers except other than those of polyolefine resin can not be used for reinforcement because they deteriorate from a shortage of heat resistant alkalinity.

On the other hand, the polyolefine resin has few hydrophilic radicals in the molecular structure and few functional groups available for theto an adhesive property ofto the cement, and therefor, the adhesion ofto the cement matrixes is extremely inferior. Accordingly, when the cement moldings reinforced with the polyolefine resin fibers are broken, the fibers are easily drawn out. Therefore, even if the impact strength and breaking energy due to drawing resistance of the fibers increase, flexural strength has not been improved greatly yet.

To improve hydrophilicity between the cement and the polyolefine resin fibers, a method in which hydrophilic high polymers such as various kinds of inorganic fine grains and polyvinyl acetate resin are added to the fibers has been proposed. However, in this case, since foreign matters are is entrapped all over the resin fibers, a drawing property is misseding and not enough fiber strength is obtained not given. In addition, nothing contributes to improving an affinity except a modifier on the surfaces of the resin fibers, and not enough reforming effects can not be obtained given compared with for the added amount added.

The pPolyolefine resin fibers comprising short fibers such as split yarn, single yarn or focusing yarn whose fineness is less than 100dt and whose fiber-length is less than 5mm produce fiber masses called "fiber balls" from their shapes. In addition, it is difficult to equally disperse the fibers in the cement because they are bulky. In <u>increasingenlarging the</u> fineness of the fibers to improve dispersibility, adhesion to the cement is inferior, and the fibers are drawn out when flexural stress is charged. Therefore, enough reinforcing effects can not be <u>obtainedgiven</u>.

In contrastOn the contrary, a method in which surfactants comprising polyoxyalkylene-alkylphenyl ether phosphate, polyoxyalkylene-fatty acid ester, and the like are applied on polypropylene resin fibers that are cut into thick filaments has been proposed (see Japanese Patent Laid-open Publication No. 116297 of 1999). Here, the filaments have irregularities having a specific average compression on fiber sections, and therein, the single yarn fineness is more than 200dt and the length is more than 5mm. However, since the surfactants have no adhesion to the polyolefine resin fibers, even if the polyolefine resin fibers adheres to the cement matrixes, enough adhesive strength can not be achieved given the rebetween. Accordingly, the cement moldings can not improve in flexural toughness.

The present invention aims to solve the above-mentioned problems. That is, it aims to provide polypropylene resin fibers for cement reinforcement which give hydrophilicity imparted to the polyolefine resin fibers, show good dispersibility and physical coupling to the cement, as well as excellent adhesion to cement matrixes, and give cement moldings improved in flexural toughness, flexural strength and impact strength[[,]]. The present invention also aims to provide fiber-reinforced molded cement made with the fiberstherewith, a method of constructing concrete structures therewith, and a method of concrete spraying therewith.

SUMMARY OF THE INVENTION

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The present invention basically provides fibers for cement reinforcement that have an increased in an index of wetting on their surface and improved in hydrophilicity. The fibers comprise specified polyolefine resin, having undergone a specified surface modification treatment on surfaces thereof. That is, the fibers are spun from polypropylene resin, and their surfaces have undergone athe surface modification treatment selected frombetween oxidation treatment and fluorination treatment so that the [[a]] fiber surfaces have an index of wetting of 38 dyn/cm or higher. Here, the index of wetting is a Japanese industrial standard, which that indicates ability for holding an adhesion between the fiber surface and the concrete. By this, an excellent hydrophilicity is given to boundary surfaces between polypropylene fibers and the cement, and cement moldings are manufactured so as to be excellent in adhesion to the cement matrixes, flexural strength and impact strength.

In this case, since the fibers are formed into monofilaments having athe single yarn fineness of 200dt or more with irregularities on their spun fiber surfaces, thea problem that contact area with the cement is decreased in generally thick fibers can be made up for to improve physical coupling to the cement. Therefore, the cement moldings excellent in flexural toughness can be manufactured.

Besides, the present invention provides polypropylene resin fibers for cement reinforcement in which corona discharge treatment is selected for oxidation treatment on their fiber surfaces. After this treatment, the fibers have an index of wetting within the range of 40-90dyn/cm.

Moreover, the present invention provides polypropylene resin fibers for cement reinforcement in which fluorinating treatment is performed on their fiber surfaces at <u>a fluorine</u> gas concentration within the range of 5-40% by volume. After this treatment, the fibers have an index of wetting within the range of 50-90dyn/cm.

Furthermore, the present invention provides cement moldings for fiber reinforcement manufactured from cement compositions to which is added an adequate amount of the above-mentioned polypropylene resin fibers in mortal mixtures including cement, fine aggregates and water.

Besides, the present invention provides a method of constructing concrete structures for manufacturing by mixing a fixed amount of the above-mentioned polypropylene resin fibers to concrete mixtures including cement, fine aggregates, coarse aggregates and water.

Moreover, the present invention provides a method of concrete spraying for spraying a mixture of mixed an adequate amount of the above-mentioned polypropylene resin fibers and to concrete mixtures including cement, fine aggregates, coarse aggregates and water on an executed surface in a fixed thickness dimension.

25 PREFERRED EMBODIMENT OF THE INVENTION

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In the present invention, the polypropylene resin used as the fiber materials is selected from among propylene homopolymer, polypropylene copolymer such as ethylene-propylene block copolymer or random copolymer, and mixtures thereof. In these, the propylene homopolymer is preferable for one for cement reinforcement, which requires high strength and heat resistance.

Especially, it is desirable to select one having an isotactic pentad fraction of 0.9 or more. Here, the isotactic pentad fraction, which is published in Macromolecules 6 925 (1973) by A. Zambelli et al, means isotactic fraction per pentad unit in a polypropylene molecule measured by using ¹³C-NMR.

The Mmelt flow rate (which is outlined in MFR) of the polypropylene resin is selected from within the range of 0.1-50g/10minutes forin view of continuous stable productivity,—[[,]].

Mmore preferably, it is selected from within the range of 1-10g/10minutes.

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Different polyolefine resins can be added to the polypropylene resin during the spinning process, if necessary. Here, as the different polyolefine resins, are listed poly (1-butene), polyethylene resin such as high density polyethylene, liner low density polyethylene, low density polyethylene, ethylene-vinyl-acetate copolymer, ethylene-alkyl-acrylate copolymer, and the like.

Antioxidants, lubricants, ultraviolet absorbents, anti-static agents, inorganic fillers, organic fillers, cross linking agents, foaming agents, nucleus agents, and the like may be mixed withto the above-mentioned polypropylene composition in accordance with using objects of the use of the composition without deviating from the present invention.

In the present invention, the polypropylene resin fiber to be spun is short one cut from monofilament having a random thickness. A manufacturing method for this is not defined specifically.[[,]] Ffor example, a manufacturing technology that extrudes filament from a round, elliptic, modified or string-shaped die is adopted.

Concretely, a publicly known melt spinning method is adopted, and spinning is carried out with a string-shaped die drawable in high magnification. In this case, polypropylene is melted and extruded from the string-shaped die, and then, a drawing treatment is performed whileas keeping the extruded string-shaped tape to form the fibers.

Although the string-shaped die has at least two nozzles connected in series, usually 5-20 nozzles are connected, and preferably 10-15 nozzles are connected.

The monofilament in the present invention may be short fibers cut from comparatively thick monofilament. In this case, in addition to the above-mentioned single layer filament, complex monofilament comprising a core layer of polypropylene component having a high melting point and a sheath layer of polypropylene component having a low melting point can be used. In this manufacturing method, polypropylene forming each layer is melted and mixed with an extruder.

Here, two discharge holes are concentrically provided on the die. The core layer comprising high melting point component is fed from the center side discharge hole of the die, and the sheath layer comprising low melting point component is extruded onto the outside thereof so as to cover the core layer to form the complex monofilament.

In this case, since essential strength depends on the materiality of the core layer, it is preferable to use propylene homopolymer, isotactic polypropylene, and the like as a high melting point component. On the other hand, as a low melting point component, it is preferable to use propylene-ethylene block copolymer, random copolymer and syndiotactic polypropylene, and the like. Such complex monofilament can prevent polypropylene resin fibers from heat deteriorations in a thermal cure at the time of concrete molding.

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Next, the monofilament has undergone <u>a</u> thermal drawing and thermal relaxing treatment. According to this, the rigidity of the filament is increased. Therefore, polypropylene monofilament suitable for cement reinforcement having a small elongation can be <u>obtainedgiven</u>. The thermal drawing is performed at a temperature <u>greater than theover a</u> softening point of polypropylene, and <u>less than thebesides under a</u> melting point thereof. Usually, the drawing temperature is within the range of 90-150°C, <u>and</u> the drawing magnification is within the range of 5-12 times, preferably within the range of 7-9 times. As a thermal drawing method, is adopted a hot roll system, a hot plate system, an infrared irradiation system, a hot air oven system, a hot water system, <u>orand</u> the like is adopted.

The Ftensile strength of the drawn polypropylene filament is 5g/dt or more, preferably 6g/dt or more. The Ftensile elongation thereof is 20% or less, preferably 15% or less. It is undesirable that the tensile strength and the tensile elongation are not respectively within these ranges because the strength of the polypropylene resin fibers for cement reinforcement becomes insufficient.

The single yarn fineness of the polypropylene monofilament is within the range of 5-10,000 decitex (dt), preferably within the range of 10dt to 6,500dt. In this case, for example, a comparatively thin one having athe single yarn fineness within the range of 5-100dt is cut in short fibers having athe fiber length within the range of 3-30mm, preferably within the range of 5-15mm.

AThe comparatively thick one having athe single yarn fineness within the range of 200-10,000dt is

cut in the fiber lengths within the range of 5-100mm, preferably within the range of 20-70mm. The fibers having athe length under 3mm come off the cement[[.]]. Aand the fibers having athe length over 100mm are inferior in the dispersibility.

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Since the fibers having athe single yarn fineness under 5dt are too thin, they easily become fiber balls because of the uniform dispersibility in the concrete mixture, thereby causing problems in execution or reinforcement. On the other hand, in case of athe single yarn fineness over 200dt, since the contacttouch area between the fibers and the concrete mixture is reduced, the fibers come to be easily drawn against the flexural stress, and the reinforcement effects are decreased. Therefore, in the present invention, for comparatively thick ones having athe single yarn fitness of 200dt or more, it is necessary to form irregularities on their surfaces as a next process of the spinning and the thermal drawing. By this, the contacttouch area between the fibers and the concrete is increased, the fibers drawing following the concrete stiffening is regulated, and the reinforcement effects can be improved. As a method for forming irregularities on their surfaces, for example, a method for embossing monofilament is listed. The embossing is performed by passing the monofilament through an embossing roller before drawing or after drawing, and irregularities are continuously formed inen a longitudinal direction of the monofilament.

Although the shape of embossing, such as the length or the depth, may be optional, it is necessary that an average compression of the fiber cross-section by crushing is within the range of 1.5/1-7/1. Here, the average compression means an average ratio of the width and the height in a cross-section of fiber having various shapes. When the average compression is under 1.5/1, there are a few irregularities on the fiber surfaces. Accordingly, there are no differences between the plane surface fibers and the reinforcing effects. On the other hand, when the average compression is over 7/1, the strength by shaping is remarkably inferior. Accordingly, the dispersibility in the concrete is apt to grow worse.

In the present invention, the above-mentioned polypropylene resin fibers have undergone a surface modification treatment selected <u>frombetween</u> oxidation treatment and fluorination treatment so that the fiber surface has an index of wetting of 38dyn/cm or more. One having <u>anthe</u> index of wetting under 38dyn/cm is undesirable because enough treatment can not be given to the polyolefine resin fibers and because <u>the</u> flexural strength and impact strength of the cement moldings can not be

improved.

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The oOxidation treatment is selected as at least a kind of treatment method from among corona discharge treatment, plasma treatment, flame-plasma treatment, electron beam irradiation treatment and ultraviolet irradiation treatment. Preferably, corona discharge treatment or plasma treatment is selected.

The corona discharge treatment is performed under a general condition, for example, the distance between the tip of electrode and the treated foundation within the range of 0.2-5.0mm. The processing amount is 5w · minute or more per 1m² polypropylene fiber, preferably within the range of 5-200w · minute/m², and more preferably within the range of 10-to 180w · minute/m². When the amount is under 5w · minute/m², the effect of the corona discharge treatment is so insufficient that the index of wetting of the fiber surface can not be set up within the above-mentioned range, and that flexural strength and impact strength of the cement moldings can not be improved.

The plasma treatment process comprises spraying an electrically neutralized excited gas on the surface of plastic base material. The excited gas is formed by eliminating charged particles from a simple substance gas such as argon, helium, krypton, neon, xenon, hydrogen, nitrogen, oxygen, ozone, carbon monoxide, carbon dioxide and sulfur dioxide or a mixed gas thereof after being electrically excited by a plasma jet. The plasma jet is formed by applying voltage between counter electrodes under an approximatelyabout atmospheric pressure and then generating plasma discharge from the mixed gas comprising, for example, nitrogen and oxygen including an oxygen concentration within the range of 5-15% by volume. In this case, the distance between the electrodes that the plastic base material passes is suitably determined by the thickness thereof, the magnitude of the applying voltage, the flow rate of the mixed gas, and the like. For example, the distance is usually within the range of 1-50mm, preferably within the range of 2-30mm. It is preferable that the voltage between the electrodes is applied so that field strength is within the range of 1-40kv/cm. Here, the frequency of AC power is within the range of 1-100kHz.

The flame plasma treatment process comprises spraying ionization plasma on the surface of the plastic base material. Here, the ionization plasma is generated in flames when natural gas or propane burns.

The electron beam irradiation treatment comprises irradiating electron beams on the surface of the plastic base material. Here, the irradiating electron beams are generated by an electron beam acceleratore. As an electron beam radiator, for example, is used a device <u>named</u> "ELECTRO CURTAIN" (name of goods) that irradiates uniform electron beams from liner filaments in the shape of a curtain.

The ultraviolet irradiation treatment process comprises irradiating ultraviolet having \underline{a} wavelength within the range of 200-400m μ on the surface of the plastic base material.

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In the fluorination treatment, for example, fluorine gas is contacted withto the polypropylene fiber surfaces in the presenceunder-existence of oxygen to form a surface-oxidizing zone thereon. According to this, the index of wetting of the surface thereof is improved in the above-mentioned range. For example, it is performed within the range of a fluorine gas concentration within the range of 5-40% by volume in the presenceunder existence of an oxygen concentration within the range of 60-95% by volume. It is preferable that the pressure is comparatively low to easily perform the reaction operation and the regulation. Specially, it is preferable that the pressure is 50 Pa or less. A type of the fluorination treatment can be selected from either a batch type or a continuous type. Treatment temperature is usually within the range of 10-100°C, preferably within the range of 10-40°C. The treatment time differs within the concentration of the fluorine gas, the pressure thereof, the treatment temperature, and the like, being usually within the range of 10 seconds to 2 hours, preferably 30-60 seconds.

In the case of the fluorination treatment by a batch process, the polypropylene resin fibers are put into a reaction container in advance, thereafter being vacuously deaerated. In addition, oxygen gas within the range of 60-95% by volume is fed therein, and then, fluorine gas within the range of 5-40% by volume is fed therein. It is desirable that the fluorination treatment is performed at the treatment temperature of 10-100°C. After the fluorination treatment, unreacting gas is eliminated from the container, and the inside thereof is efficiently substituted and ventilated by inactive gas to produce fluorination-treated polypropylene fibers.

Polypropylene fibers for reinforcement cement of the present invention can be used in various embodiments with mixed in mortars or concretes.

For example, are listed a method for manufacturing cement moldings by forming

therewith, a method for constructing concrete structures or mortar structures by placing and painting therewith, a method for constructing concrete structures or mortar structures by spraying therewith, and the like.

When the polypropylene fibers are mixed in mortars, they are used with mixed in cement, fine aggregates, water and a proper amount of admixture.

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Besides, when they are mixed in concrete, they are used with mixed in cement, fine aggregates, coarse aggregates, water and a proper amount of admixture. Here, as the cement, are listed hydraulic cement such as Portland cement, blast furnace cement, silica cement, fly ash cement, white Portland cement, alumina cement, and the like, and non-hydraulic cement such as gypsum and lime. As the fine aggregates, are listed river sand, sea sand, pit sand, silica sand, glass sand, iron sand, ash sand, another artificial sand, and the like. As the coarse aggregates[[,]] are listed gravel, ballast, crushed stone, slag, various artificial lightweight aggregates, and the like.

In manufacturing the cement moldings, as a method for mixing polypropylene fibers in cement, can be used publicly known methods such as a method for dispersing them in cement powder, a premix method for dispersing them in cement slurry, and a spray-up method for spraying cement, polypropylene fibers and water at the same time, and the like.

The composite amount of the polypropylene fibers is 0.1-10% by weight to the cement, preferably within the range of 0.5-5% by weight. AThe composite amount of 0.1% by weight or less is undesirable because reinforcement effects are inferior. AThe composite of 10% by weight or more is undesirable because flexural strength is inferior and uniform dispersion is difficult.

The Tthus given cement slurry is molded by a publicly known method such as a watermark molding method, an extrusion molding method, an injection molding method according to purposes. Such a molding is cured by a natural curing method byfor being left in the atmosphere or in water at an ordinary temperature for about 10 days or by an autoclave method byfor being disposed at a temperature of 100-200°C after being left at the ordinary temperature for a few days to produce a cement molding.

The purposes of the cement moldings in the present invention cover all sorts of cement products. For example, they are used in wall materials, floor material concretes, finishing mortars, waterproofed concretes, slate roof materials <u>for</u>as buildings, or pavement materials such as roads and

airstrips, road materials such as road-signs and gutters, pipes such as sewer pipes and cable ducts, fish banks, bulkhead blocks, tetrapots, and various organizations such as sleepers, benches and flower pots.

In using the polypropylene fibers to execute mortar structures, they are mixed with cement, fine aggregates, water and a proper amount of admixture at the same time or in the state that the mortars are tempered up, and placed and painted. Besides, in executing concrete structures, the polypropylene fibers are mixed with cement, fine aggregates, water and a proper amount of admixture at the same time or in the state that the concretes are tempered up, and placed and painted.

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In the present invention, it is important to mix and disperse the polypropylene fibers of 4-19kg, preferably 6-14kg, to 1m³ of concrete mixtures comprising cement, fine aggregates, rough aggregates, water and the like. When the mixed amount of the polypropylene fibers is over 19kg, flexural toughness does not increase because the fibers do not uniformly disperse in the concrete. On the other hand, in the mixed amount of less than 4 kg, bounces at spraying are large and reinforcement effects after curing are small.

In <u>a</u> mixing method <u>according to the present invention of this case</u>, it is preferable that the concrete mixtures comprising cement, fine aggregates, rough aggregates, water and the like are input to form base concrete and the polypropylene fibers are input and mixed therein after mixing the base concrete. Although mixing time depends on a mixed amount per a time, it is proper that the mixing time is for 45-90 seconds in the base concrete and for 45-90 seconds after inputting the polypropylene fibers.

In addition, in a method of spray concreting, it is preferable that slump is adjusted within the range of 8-21cm in using the polypropylene fibers in the above-mentioned composite amount. A The slump under 8cm is undesirable because spraying work is difficult, and athe slump over 21cm is undesirable because the bounces become large. When the slump is within the above-mentioned range, it is effective in a spray nozzle to be perpendicular to athe sprayed surface with and to make the distance therebetween within the range of 0.5-1.5m.

The Tthus produced concrete mixture is used to cover tunnels (including batter piles and shafts) and large cavity structures, to prevent slop surfaces, slant surfaces or wall surfaces from weathering, peeling and flaking, and to mend and reinforce tunnels and bridges as a shotcrete.

The following examples will explain about <u>the efficiency</u> of the polypropylene fibers in the invention.

Example 1:

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Polypropylene resin (MFR = 1.0g/10 minutes) was fed into an extruder, extruded from string-shaped nozzles having $2mm \phi \times 10$ bores at 230° C resin temperature, and drawn by a hot plate contact type drawing method. A drawing temperature was 130° C, an annealing temperature was 135° C, and a drawing magnification was 12 times. The given drawn yarn had the single yarn fineness of 50dt.

The drawn yarn surface had undergone a corona discharge treatment as a surface oxidation treatment under 20 w·minute per 1m². The given polypropylene resin drawn yarn surface had an index of wetting of 42 dyn/cm.

The polypropylene resin drawn yarn was cut in 10mm length to form short fibers.

Molding for cement moldings was based on JISR5201. That is, 100 wt. parts Portland cements and 200 wt. parts standard sands were mixed sufficiently, and therein, the fibers of 5 wt. parts and water of 65 wt. parts are added. Then they were mixed uniformly on the whole. Thereafter, they were poured into a mould of $40 \text{mm} \times 40 \text{mm} \times 160 \text{mm}$, left in the atmosphere for 48 hours at the ordinal temperature, thereafter cured in the autoclave for 20 hours at 165°C .

The given molding had a flexural strength of 26.0 MPa and Charpy impact strength of 9.5KJ/m². The dispersibility was good.

(Test Method)

- (1) MFR: Based on JISK6922-1
- (2) Flexural strength: Based on JISA1408
- (3) Charpy impact strength: Based on JISB7722
- (4) Dispersibility Appreciation: The surface of cement slurry comprising polypropylene fibers and cement was appreciated by visual observation.

Example 2:

The same treatments as Example 1 except the conditions that the polypropylene resin

drawn yarn surface had undergone the corona discharge treatment on at the rate of 30w·minute per 1m² and the index of wetting the surface was 45 dyn/cm were performed.

The given moldings had the flexural strength of 26.5 MPa, Charpy impact strength of 9.8 KJ/m². The dispersibility was good.

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Example 3:

A drawn yarn was produced similarly with Example 1 and cut in 10mm length to form short fibers. The short fibers were put into a reaction container, and thereafter, the container was placed under amade in vacuum. Gaseous oxygen of 80% by volume and gaseous fluorine of 20% by volume were injected in order, and these were reacted at 20°C under 10 Pa pressure. The given fibers had the surfaces having the index of wetting of 60 dyn/cm.

Cement moldings were molded similarly with Example 1 by using these short fibers. The given moldings had athe flexural strength of 28.0 MPa and Charpy impact strength of 10.5 KJ/m². The dispersibility was good.

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Comparative example 1:

As a finishing agent, was produced a finishing agent liquid mixed 50% by weight polyoxyethylene nonyl phenyl ether phosphate (HLB = 8.0) and 50% by weight polyoxyethylene oleic acid ester (HLB = 9.0) was produced. The fibers were dipped therein, and thereafter dried to be applied the finishing agent of 1% by weight. The treatments except these were the same as in Example 1.

The given moldings had <u>athe</u> flexural strength of 19.0 MPa and <u>a Charpy</u> impact strength of 6.5 KJ/m². The dispersibility of the fibers was good.

25 Comparative example 2:

As the finishing agent, was produced a finishing agent liquid mixed 70% by weight polyoxyethylene nonyl phenyl ether phosphate (HLB = 8.0) and 30% by weight polyoxyethylene oleic acid ester (HLB = 9.0) was produced. The fibers were dipped therein, and thereafter dried to be applied the finishing agent of 1% by weight. The treatments except these were the same as in

Example 1.

The given moldings had <u>athe</u> flexural strength of 16.5 MPa and <u>a</u> Charpy impact strength of 3.5 KJ/m². The dispersibility of the fibers was no good.

5 Comparative example 3:

As the finishing agent, was produced a finishing agent liquid mixed 30% by weight polyoxyethylene nonyl phenyl ether phosphate (HLB = 8.0) and 70% by weight polyoxyethylene oleic acid ester (HLB = 9.0) was produced. The fibers were dipped therein, and thereafter dried to be applied the finishing agent of 1% by weight.

The given moldings had <u>athe</u> flexural strength of 17.5 MPa and <u>a Charpy</u> impact strength of 2.8 KJ/m². The dispersibility of the fibers was approximately no good.

Example 4:

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(1) Producing fibers

Polypropylene resin (MFR = 4.0 g/10 minutes, Tm = 163°C) was input into an extruder, spun from circular nozzles and cooled. Thereafter, it was drawn at 115°C hot drawing temperature, 120°C hot relaxation temperature and 7-8 times drawing magnifications in accordance with a hot air oven type drawing method to produce [[a]] several kinds of monofilament having different fineness. Then, the monofilament wasis formed into polypropylene monofilament having irregularities on the surface different in an average compression by using a slanted grid emboss roll and hard rubber roll and by changing emboss nip pressure.

As the surface oxidation treatment, a corona discharge treatment was performed on the polypropylene monofilament surface at 30w·minute per 1m². The given monofilament had <u>athe</u> surface <u>with anof the</u> index of wetting of 45 dyn/cm. The polypropylene monofilament was cut in 30mm length to be formed polypropylene fibers.

(2) Appreciation Test

The given polypropylene resin fibers were tested <u>for theirabout a reinforcement effect of</u> concrete by the following methods. The results are shown in table 1.

① Materials and Compounding ratio

Cement: $430 \text{ kg/m}^3 \text{ high early strength Portland cement (specific density = 3.12)}$

Fine aggregate: 1123 kg/m³ pit sand made in Kisarazu (specific gravity in saturated surface-dry condition = 2.60)

Rough aggregate: 491 kg/m³ crushed stone made in Oumi 1505 (specific gravity in saturated surface-dry condition = 2.65)

Water: 215 kg/m³ service water

Fiber: 1% as volume

② A method for mixing concrete

Using a forced pan mixer having <u>a</u> mixing volume of 100 I, 1 batch was conducted at 60 I.

A temperature at the time when the concrete was tempered was approximately 20°C. The mixing method was conducted by inputting fine aggregate, cement, water and rough aggregate, mixing them for 45 seconds, thereafter adding reinforcement fibers <u>whileas</u> rotating the mixer, mixing these for 60 seconds, and discharging.

③ Producing test pieces

This was based on Civil Engineering Standard "Strength Of Steel Fiber Reinforcement Concrete And How To Make Test Pieces For Compressed Toughness" (JSCE F552-1983). The test pieces were taken off from the mold after 24 hours, and a water curing was conducted till material ages of 7 days.

4 A method for testing

This was based on Civil Engineering Standard "Compressed Strength Of Steel Fiber Reinforcement Concrete And How To Make Test Pieces For Compressed Toughness" (JSCE F551-1983) and Civil Engineering Standard "Flexural Strength Of Steel Fiber Reinforcement Concrete And How To Make Test Pieces For Compressed Toughness" (JSCE F552-1983).

Example 5 and Example 6:

They were conducted similarly with Example 4 except <u>thatehanging</u> the fineness and the compression of polypropylene resin fiber <u>were changed</u>. The results are shown in table 1.

Example 7:

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Polypropylene monofilament was produced similarly with Example 4, and cut in short fibers having 30mm length.

After the short fibers were put into a reaction container, the container was placed under was placed under was under was oxygen of 80% by volume and gaseous fluorine of 20% by volume were lifed into the container in order, and a reaction was performed at 20°C under the pressure of 10 Pa. The given polypropylene short fibers had the surfaces having anthe index of wetting of 60 dyn/cm.

The polypropylene short fibers were tested <u>for theirabout a</u> concrete reinforcement effect similarly with Example 4. The result is shown in table 1.

10 Example 8 and Example 9:

They were conducted similarly with Example 7 except <u>for</u> changing the fineness and the compression of polypropylene resin fiber. The results are shown in table 1.

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Table 1

	Fiber	Fineness	Compression	Weight	Flexural toughness	Compressed strength
	(-)	(dt)	(-)	(kg/m³)	(kgf · cm)	(N/mm ²)
Example 4	PP	3000	4.2/1	9.2	425	38.1
Example 5	PP	6000	6.4/1	9.2	430	38.3
Example 6	PP	500	2.6/1	9.2	418	37.8
Example 7	PP	3000	4.2/1	9.2	451	38.1
Example 8	PP	6000	6.4/1	9.2	461	38.3
Example 9	PP	500	2.6/1	9.2	442	37.8
Comparative example 4	PP	3000	4.2/1	9.2	317	37.5
Comparative example 5	PP	6000	6.4/1	9.2	325	37.8
Comparative example 6	PP	500	2.6/1	9.2	310	37.3
Comparative example 7	steel	φ 0.6 mm	3.0/1	78.0	330	37.5
Comparative example 8	PVA	4000	1.4/1	13.0	151	35.7

CLAIMS

1. Polypropylene fiber for cement reinforcement, comprising:

fibers spun out of polypropylene resin, said fibers having undergone a surface modification treatment selected between oxidation treatment and fluorination treatment so that a fiber surface has an index of wetting of 38 dyn/cm or more.

2. Polypropylene fibers for cement reinforcement, comprising:

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monofilament spun out of polypropylene resin, said monofilament having single yarn fineness of 200 dt or more with irregularities on their surfaces, and having undergone a surface modification treatment selected between oxidation treatment and fluorination treatment so that a monofilament surface has an index of wetting of 38 dyn/cm or more.

3. Polypropylene fibers for cement reinforcement of claim 1 or claim 2,

wherein the oxidation treatment is selected between corona discharge treatment and plasma treatment so that a fiber surface has an index of wetting within the range of 40 to 90 dyn/cm after the treatment.

4. Polypropylene fibers for cement reinforcement of claim 1 or claim 2,

wherein the fluorination treatment is conducted under fluorine gas in a concentration within the range of 5 to 40 % by volume so that a fiber surface has an index of wetting within the range of 50 to 90 dyn/cm after the treatment.

5. Molded cement, comprising:

cement composition having an adequate amount of polypropylene fiber of claim 1 added to mortar mixture including cement, fine aggregate and water.

6. Method of constructing concrete structure which comprises mixing a fixed amount of polypropylene fiber of claim 2 to concrete mixture including cement, fine aggregate, rough aggregate and water.

7. Method of spray concreting which comprises mixing an adequate amount of polypropylene fiber of claim 2 to concrete mixture including cement, fine aggregate, rough aggregate and water, spraying the mixture on a formation surface in a fixed thickness.

ABSTRACT

Polyolefine resin fibers for cement reinforcement which have hydrophilicity impareted thereto and show excellent adhesion to cement matrices. The fibers are ones which have been formed from a polypropylene resin by spinning and have undergone a surface modification treatment selected frombetween oxidation treatment and fluorination treatment so that the fiber surface has an index of wetting of 38 dyn/cm or higher. The polypropylene resin fibers for cement reinforcement, when used in the application or placing of various mortars or concretes, give cement moldings improved in flexural toughness, such as flexural strength and impact strength.